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DURABLE MEDICAL EQUIPMENT

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WHEELCHAIRS FOR NURSING HOME PARTICIPANTS

MO HealthNet Division (MHD) will reimburse for medically necessary custom wheelchairs for participants residing in a nursing facility. A custom wheelchair is defined as follows:

A. Any wheelchair with a custom seating system. A custom seating system is a wheelchair seating system which is individually made for a patient using a plaster model of a patient, a computer generated model of the patient (i.e. CAD-CAM technology), or the detailed measurements of the patient to create either: (a) a molded, contoured, or carved (foam or other suitable material) custom-fabricated seating system that is incorporated into the wheelchair base; or (b) a custom seating system made from multiple pre-fabricated components or a combination of custom fabricated materials and pre-fabricated components which have been configured and attached to the wheelchair base or incorporated into a wheelchair seat and/or back in a manner that the wheelchair could not be easily re-adapted for use by another individual.

To qualify for a custom seating system, an individual must meet all the requirements of a custom fabricated seat cushion or a custom fabricated back cushion as described in Section 13.29.G of the Durable Medical Equipment Provider Manual. The prior authorization request must document the following:

- 1. Why a prefabricated system is not sufficient to meet the participant's seating and positioning needs.
- 2. What orthopedic deformity is present and it's fixed or flexible presentation.
- 3. What altered muscle tone is present and its increased or decreased presentation that affects seating and positioning.
- 4. Why any existing system is not meeting the participant's seating and positioning needs.

- B. A specially sized or constructed wheelchair that is provided to a participant whose anatomical measurements require the following:
- 1. A wheelchair seat width of 25 inches or more; or
- 2. A wheelchair with a weight capacity for 400 or more pounds; or
- 3. A wheelchair with a seat to floor height of less than 15 ½ inches
- C. A wheelchair with a manual tilt feature for a participant who has absent or impaired sensation in the area of contact with the seating surface or inability to carry out a functional weight shift due to one of the following diagnoses:
- 1. spinal cord injury resulting in quadriplegia or paraplegia (344.00-344.1)
- 2. other spinal cord diseases (336.0-336.3)
- 3. multiple sclerosis (340)
- other demyelinating disease (341.0-341.9);
- 5. cerebral palsy (343.0-343.9);
- 6. anterior horn cell diseases including amyotrophic lateral sclerosis (335.0-335.21, 335-23-335.9)
- 7. post polio paralysis (138)
- 8. traumatic brain injury resulting in quadriplegia (344.09);
- 9. spina bifida (741.00-741.93)
- 10. childhood cerebral degeneration (330.0-330.9)
- 11. Alzheimer's disease (331.0)
- 12. Parkinson's disease (332.0)
- 13. current stage II or greater pressure ulcer (707.03, 707.04, 707.05) or past history of a stage II or greater pressure ulcer (707.03, 707.04, 707.05) on the area of contact with the seating surface (trunk, spine or pelvis)

PRIOR AUTHORIZATION REQUESTS FOR WHEELCHAIRS

When submitting a prior authorization request for a custom wheelchair or power mobility device, there must be comprehensive written documentation submitted with the prior authorization request that clearly and specifically explains all the following:

- The diagnosis/comorbidities and conditions relating to the need for a custom or power wheelchair
- Description and history of limitations/functional deficits
- Description of physical and cognitive abilities to utilize equipment
- History of previous interventions/past use of mobility devices
- Description of existing equipment, age and specifically why it is not meeting the participants needs
- Why a less costly mobility device is unable to meet the participants needs (i.e., cane, walker, standard wheelchair)
- Documentation and justification of medical necessity of recommended mobility device, accessories and positioning components
- Documentation/explanation of participant's ability to safely tolerate/utilize the recommended equipment

Provider Bulletins are available on the MO HealthNet Division (MHD) (Formerly the Division of Medical Services) Web site at http://dss.mo.gov/mhd/providers/pages/bulletins.htm. Bulletins will remain on the Provider Bulletins page only until incorporated into the provider manuals as appropriate, then moved to the Archived Bulletin site.

MO HealthNet News: Providers and other interested parties are urged to go to the MHD Website at http://dss.missouri.gov/mhd/global/pages/mednewssubscribe.htm to subscribe to the electronic mailing list to receive automatic notifications of provider bulletins, provider manual updates, and other official MO HealthNet communications via e-mail.

MO HealthNet Managed Care: The information contained in this bulletin applies to coverage for:

- MO HealthNet Fee-for-Service
- Services not included in MO HealthNet Managed Care

Questions regarding MO HealthNet Managed Care benefits should be directed to the patient's MO HealthNet Managed Care health plan. Before delivering a service, please check the patient's eligibility status by swiping the red MO HealthNet card or by calling the Interactive Voice Response (IVR) System at 573-635-8908 and using Option One for the red or white card.

Provider Communications Hotline 573-751-2896